SUPPLEMENTAL INFORMATION DISCLOSURE STATEMENT LIST

(Use as many sheets as necessary)

Complete if Known	
Application Number	10/579,149
Filing Date	January 19, 2007
First Named Inventor	Kortney L. Klinkel
Group Art Unit	1611
Examiner Name	Joseph S. Kudla

U.S. PATENT DOCUMENTS

Examiner's Initials	Cite No.	Document No.	Date	Name	Class	Subclass	Filing Date (if appropriate
	B1	3,279,918	10-18-1966	Gevaert Photo-Production N.V.	90	1	
	B2	3,297,710	1-10-1967	E.I. du Pont de Nemours & Co.	260	309	
	B3	4,089,747	5-16-1978	Eastman Kodak Co.	435	10	
	B4	4,423,046	12-27-1983	Sterling Drug Inc.	514	228.8	
	B5	4,466,976	8-21-1984	Schering Aktiengesellschacft	514	397	
	B6	4,585,771	4-29-1986	Schering Aktiengesellschacft	514	220	
	B7	4,705,855	11-10-1987	Rottapharm SpA	544	370	
	B8	4,721,670	1-26-1988	Fuji Photo Film Co.	435	28	
	B9	4,902,705	2-20-1990	UBE Industries, Ltd. et al.	514	397	
	B10	4,970,226	11-13-1990	Harbor Branch Oceanographic Institution, Inc.	514	397	
	B11	5,024,935	6-18-1991	Eastman Kodak Co.	435	7.1	
	B12	5,047,318	9-10-1991	Eastman Kodak Co.	435	5	
	B13	5,496,702	3-5-1996	Johnson & Johnson Chemical Diagnostics, Inc.	435	7.9	
	B14	5,514,550	5-7-1996	Johnson & Johnson Chemical Diagnostics, Inc.	435	6	
	B15	5,656,644	8-12-1997	SmithKline Beecham Corp.	514	341	
	B16	5,686,455	11-11-1997	SmithKline Beecham Corp.	514	256	
	B17	5,693,589	12-2-1997	Eastman Kodak Co.	503	227	
	B18	5,753,687	5-19-1998	Ontogen Corp.	514	396	
	B19	5,916,891	6-29-1999	SmithKline Beecham Corp.	514	256	
	B20	5,945,418	8-31-1999	Vertex Pharmaceuticals, Incorp.	514	248	
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	B22	6,117,609	9-12-2000	Brother Kogyu Kabushiki Kaisha	430	138	

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	B23	6,194,441	2-27-2001	Zeneca Ltd.	514	340	
	B24	6,268,370	7-31-2001	SmithKline Beecham Corp.	514	256	
	B25	6,521,655	2-18-2003	Ortho-McNeill Pharmaceuticals, Inc.	514	397	
	B26	2007/0123553	5-31-2007	Lorus Therapeutics Inc.	514	285	
	B27	2004/0127527	7-1-2004	Mitsuya et al.	514	365	
	B28	2004/0176601	9-9-2004	Goulet et al.	546	46	
	B29	2007/0105929	5-10-1997	Al-Qawasmeh	514	393	

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	B30	CA 2,351,694	7-22-1993	SmithKline Beecham Corp.	
	B31	EP 165588	12-27-1985	Fuji Photo Film Co. Ltd.	
	B32	JP 2002-275458	9-25-2002	Fukuoka Prefecture	Abstract only
	B33	JP 2002-364578	12-18-2002	Hitachi Ltd.	Abstract only
	B34	WO 1995/03297	2-2-1995	SmithKline Beecham Corp.	
	B35	WO 1996/18626	6-20-1996	F. Hoffman-La Roche AG Harmon	
	B36	WO 1998/27108	6-25-1998	Fujisawa Pharmaceutical Co. Ltd.	
	B37	WO 1999/01128	1-14-1999	Neurogen Corp.	
	~~B38~~	~~WO 2000/33836~~	~~6-15-2000~~	~~Ontogen Corp.~~	
	B39	WO 2003/004023	1-16-2003	Schering Aktiengesellschacft	
	B40	WO 1993/014081	7-22-1993	SmithKline Beecham Corp.	
	B41	WO 1998/027065	6-25-1998	Ontogen Corp.	
	B42	WO 2000/078761	12-28-2000	Sepracor Inc.	
	B43	WO 2002/046168	6-13-2002	Astrazeneca AB	
	B44	WO 2004/016086	2-26-2004	Lorus Therapeutics Inc.	
	B45	WO 2005/047266	11-15-2004	Lorus Therapeutics Inc.	

NON-PATENT DOCUMENTS

Examiner's Initials	Cite No.	Non-Patent Citations (include Author, Title, Publisher, Relevant Pages, Date and Place of Publication)
	B46	Abdel-Meguid *et al.*, "An orally bioavailable HIV-1 protease inhibitor containing an imidazole-derived peptide bond replacement: Crystallographic and pharmacokinetic analysis," *Chemistry*, 1994, 33:11671-11677
	B47	Adams *et al.*, "Pyrimidinylimidazole inhibitors of p38: Cyclic N-1 imidazole substituents enhance p38 kinase inhibition and oral activity," *Bioorg. Med. Chem. Lett.*, 2001, 11:2867-2870

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	B48	Antolini *et al.*, "Analogues of 4,5-bis(3,5-dichlorophenyl)-2-trifluoromethyl-1H-imidazole as potential antibacterial agents," *Bioorg. Med. Chem. Lett.*, 1999, 9:1023-1028
	B49	Armesto *et al.*, "A new site selective synthesis of benzoin esters, synthesis of symmetrically and unsymmetrically substituted benzils," *Synthesis*, 1988, 799-801
	B50	Bhaduri *et al.*, "Potential Antifertility Agents. Synthesis of 2,4,5-Substituted Imidazoles", Central Drug Res. Inst. Lucknow, India, *Indian J. Chem.*, 1966, 4(9):419-420
	B51	Botana *et al.*, "*p-(1H-Phenanthro[9,10-d]imidazol-2-yl)- Substituted Calix[4]arene, a Deep Cavity for Guest Inclusion*", Departmento de Quimica Organica, Universidad Autonoma de Madrid, Spain, *Organic Letters*, 2004, 6(7):1091-1094
	B52	Bu *et al.*, "A novel approach to synthesis of tricyanovinylthiophene for heterocyclic imidazole nonlinear optical chromophores," *Tetrahedron Lett.*, 1996, 37:7331-7334
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	B56	Cuenda, *et al.*, "SB 203580 is a specific inhibitor of a MAP kinase homologue which is stimulated by cellular stresses and interleukin-1," *Febs Lett*, 1995, 364:229-33
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	B58	Demirayak *et al.*, "Synthesis of Certain Derivatives of Ethyl α-[(phenanthro[9,10-d]imidazol-2-yl)phenoxy]alkanoate", *Acta Pharmaceutica Turcica*, 1989, 31(1):19-25
	B59	Downey *et al.*, "Degradation of DNA by 1,-10-phenanthroline," *Biochem Biophys Res Commun*, 1980, 93(1):264-70
	B60	Fischer *et al.*, "Dissociation constants of the conjugate acids of substituted benzyl phenyl ketones and of alkyl-substituted benzophenomes," *J. Am. Chem. Soc.* 1961, 83:4208-4210
	B61	Gales *et al.*, "Characterization of pseudomonas aeruginosa isolates: Occurense rates, antimicrobial susceptibility patterns, and molecular typing in the global SENTRY antimicrobial surveillance program, 1997-1999," *Clin. Infect. Dis.*, 2001, 32:S146-155
	B62	Guijarro *et al.*, "The reaction of active zinc with organic bromides," *J. Am. Chem. Soc.*, 1999, 121:4155-4157
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	B65	Krieg *et al., Naturforsch.* 1967, 22b:132-141 (English translation)
	B66	Kimura *et al.*, "*Preparation of 4-(4,5-diphenyl-1H-imidazol-2-yl)benzaldehyde and its Practical Synthetic Use in the Synthesis of Unsymmetrically Substituted Imidazoles*", Department of Chemistry, Okayama University, Okayama Japan, ITE Letters on Batteries, New Technologies & Medicine, 2002, 3(1), pp. 30-34
	B67	Lee *et al.*, "A protein kinase involved in the regulation of inflammatory cytokine biosynthesis," *Nature*, 1994, 327:739-745
	B68	Lewis J.R., "Muscarine, imidazole, oxazole, thiazole and peptide alkaloids, and other miscellaneous alkaloids," *Nat. Prod. Rep.*, 1998, 15:371-395
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	B72	LoGrasso *et al.*, "Kinetic mechanism fro p38 MAP kinase," *Biochemistry*, 1997, 36:10422-10427
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	B75	Moylan *et al.*, "*Synthesis and Nonlinear Optical Properties of Donor-Acceptor Substituted Triaryl Azole Derivatives*", Almaden Res. Cent. IBM, San Jose, CA, USA, Chemistry of Materials, 1993, 5(10):1499-1508
	B76	Pan *et al.*, "DNA-binding proteins as site-specific nucleases," *Mol Microbiol*, 1994, 12(3):335-42
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	B78	Sarshar *et al.*, "Imidazole libraries on solid support," *Tetrahedron Lett.* 1996, 37:835-838
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	B84	Zeytinoglu *et al.*, "*Mutagenicity Assay in Salmonella for Thirteen 2-Substituted-1H-phenanthro (9,10-d) Imidazoles*", Department of Biology, Faculty of Science, Anadolu University, Eskisehir, Turkey, Drug and Chemical Toxicology, 26(4):245-257
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